

## Product datasheet for PH319727

### COG8 (NM\_032382) Human Mass Spec Standard

#### Product data:

Product Type:	Mass Spec Standards
Description:	COG8 MS Standard C13 and N15-labeled recombinant protein (NP_115758)
Species:	Human
Expression Host:	HEK293
Expression cDNA Clone or AA Sequence:	RC219727
Predicted MW:	68.4 kDa
Protein Sequence:	>RC219727 protein sequence Red=Cloning site Green=Tags(s)

MATAATIPSVATATAAALGEVEDEGLLASLFRDRFPEAQWRERPDVGRYLRELSGGLERLRREPERLAE  
ERAQLLQQTRDLAFANYKTFIRGAECTERIHRLFQDVEASLGRLLDRLPSPFQQSCRNFVKEAEEISSNRR  
MNSLTLNRHTEILEILEIPQLMDTCVRNSYYEEALELAAYVRRLERKYSSIPVIQIGIVNEVRQSMQLMLS  
QLIQQLRTNIQLPACLRVIQYLRRMDVFTEALRVKFLQARDAWLSILTAIPNDPPYFHITKTIEASRV  
HLFDIITQYRAIFSDPPLPPAMGEHTVNESAI FHGWVLQKVSQFLQVLETDLYRIGGHLDSLGLGQCM  
YFGLSFSRVGADFRGQLAPVFQVAISTFQKAIQETVEKFQEEEMNSYMLISAPAILGTSNMPAAVPATQP  
GTLQPPMVLLDFPPLACFLNNILVAFNDLRLCCPVALAQDVTGALEDALAKVTKIILAFHRAEEAAFSSG  
EQELFVQFCTVFLEDLVPYLNRLQVLFPPAQIAQTLGIPPTQLSKYGNLGHVNIQAIQEPLAFILPKRE  
TLFTLDDQALGPELTAPAPEPPAEPRLEPAGPACPEGGRAETQAEPSPVGP

TRTRPLEQKLI SEEDLAANDILDYKDDDDKV

Tag:	C-Myc/DDK
Purity:	> 80% as determined by SDS-PAGE and Coomassie blue staining
Concentration:	>0.05 µg/µL as determined by microplate BCA method
Labeling Method:	Labeled with [U- 13C6, 15N4]-L-Arginine and [U- 13C6, 15N2]-L-Lysine
Buffer:	25 mM Tris-HCl, 100 mM glycine, pH 7.3
Storage:	Store at -80°C. Avoid repeated freeze-thaw cycles.
Stability:	Stable for 3 months from receipt of products under proper storage and handling conditions.
RefSeq:	<u><a href="#">NP_115758</a></u>
RefSeq Size:	2522
RefSeq ORF:	1836



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**Synonyms:** CDG2H; DOR1  
**Locus ID:** 84342  
**UniProt ID:** [Q96MW5](#), [A0A024R6Z6](#)  
**Cytogenetics:** 16q22.1

**Summary:** This gene encodes a protein that is a component of the conserved oligomeric Golgi (COG) complex, a multiprotein complex that plays a structural role in the Golgi apparatus, and is involved in intracellular membrane trafficking and glycoprotein modification. Mutations in this gene cause congenital disorder of glycosylation, type IIh, a disease that is characterized by under-glycosylated serum proteins, and whose symptoms include severe psychomotor retardation, failure to thrive, seizures, and dairy and wheat product intolerance. [provided by RefSeq, Jul 2008]

**Protein Families:** Druggable Genome

### Product images:



Coomassie blue staining of purified COG8 protein (Cat# [TP319727]). The protein was produced from HEK293T cells transfected with COG8 cDNA clone (Cat# [RC219727]) using MegaTran 2.0 (Cat# [TT210002]).